

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁵ : C08J 9/08, 9/10, 9/12	A1	(11) International Publication Number: WO 93/15132 (43) International Publication Date: 5 August 1993 (05.08.93)
(21) International Application Number: PCT/US92/10932 (22) International Filing Date: 17 December 1992 (17.12.92) (30) Priority data: 07/831,123 4 February 1992 (04.02.92) US (71) Applicant: THE DOW CHEMICAL COMPANY [US/ US]; 2030 Dow Center, Abbott Road, Midland, MI 48640 (US). (72) Inventor: PARK, Chung, P. ; 8762 Stoneridge Court, Pick- erington, OH 43147 (US). (74) Agent: DEAN, J., Robert., Jr.; The Dow Chemical Com- pany, Patent Department, P.O. Box 515, Granville, OH 43023-0515 (US).		(81) Designated States: CA, JP, NO, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: DIMENSIONALLY-STABLE POLYPROPYLENE FOAM EXPANDED WITH INORGANIC BLOWING AGENTS (57) Abstract Disclosed is a process for making low density, dimensionally-stable, extruded propylene polymer foam comprised primarily of an expanded propylene polymer material wherein the foam has a blowing agent comprising at least 15 percent by weight of one or more inorganic blowing agents. The foam has a density of from 10 to 150 kilograms per cubic meter, and an average cell wall thickness of less than 35 micrometers. Further disclosed is a foam obtainable from the process.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
BB	Barbados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BG	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	IE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SK	Slovak Republic
CI	Côte d'Ivoire	LJ	Liechtenstein	SN	Senegal
CM	Cameroon	LK	Sri Lanka	SU	Soviet Union
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	MC	Monaco	TG	Togo
DE	Germany	MG	Madagascar	UA	Ukraine
DK	Denmark	ML	Mali	US	United States of America
ES	Spain	MN	Mongolia	VN	Viet Nam
FI	Finland				

DIMENSIONALLY-STABLE POLYPROPYLENE FOAM EXPANDED
WITH INORGANIC BLOWING AGENTS

5 This invention relates to a low density,
dimensionally-stable, extruded propylene polymer foam
blown with a blowing agent comprising partly or entirely
one or more inorganic blowing agents. The foam offers
excellent cushioning properties.

10 Propylene polymer foams are known in the art,
and have been found useful in many applications such as
seen in U.S. Patent No. 3,481,455, 3,871,897, 3,932,569,
4,522,955, and 4,832,770. Such foams have been made
15 with a wide variety of organic and inorganic blowing
agents.

20 Given current environmental concerns over the
use of organic blowing agents, it would be desirable to
have an extruded propylene polymer foam blown partly or
solely with inorganic blowing agents. It would be
further desirable to have such a dimensionally-stable,
low density foam (for example, 10 to 70 kilograms per
cubic meter) for use in cushioning applications. It
25 would also be desirable to have such a foam that is

partially open-cell (for example, at least 5 percent) since open-cell foams are easier to produce and dimensionally more stable than corresponding closed-cell foams.

5 According to the present invention, there is a low density, dimensionally-stable, extruded propylene foam comprising a propylene polymer material wherein the foam has a blowing agent comprising at least 15 percent
10 by weight of one or more inorganic blowing agents. The propylene polymer material comprises at greater than 50 percent by weight propylene monomeric units. The foam has a density of from 10 to 150 kilograms per cubic meter. Foam preferably has a density of from 10 to 70
15 kilograms per cubic meter.

 Further according to the present invention, there is a process for making a dimensionally-stable, extruded propylene polymer foam. The process comprises
20 the steps of (a) heating the propylene polymer material comprising greater than 50 weight percent propylene monomeric units to form a melt polymer material; (b) incorporating into the melt polymer material a blowing agent comprising at least 15 weight percent of one or
25 more inorganic blowing agents to form a foamable gel; (c) extruding the foamable gel through a die to form the foam.

 Suitable propylene polymer materials include
30 propylene homopolymers (polypropylene) and copolymers of propylene and copolymerizable ethylenically unsaturated comonomers. The propylene polymer material may further include non-propylenic polymers. The propylene polymer material may be comprised solely of one or more propylene homopolymers, one or more propylene

5 copolymers, a blend of one or more of each of propylene homopolymers and copolymers, or blends of any of the foregoing with a non-propylenic polymer. Regardless of composition, the propylene polymer material comprises at greater than 50 and preferably at least 80 weight percent of propylene monomeric units.

10 Suitable monoethylenically unsaturated comonomers include olefins, vinylacetate, methylacrylate, ethylacrylate, methyl methacrylate, acrylic acid, itaconic acid, maleic acid, maleic anhydride. The propylene copolymer preferably comprises 20 percent or less by weight of the ethylenically unsaturated comonomer.

15 Suitable non-propylenic polymers incorporatable in the propylene polymer material include high, medium, low, and linear density polyethylenes, polybutene-1, ethylene-acrylic acid copolymer, ethylene-vinyl acetate copolymer, ethylene-propylene rubber, styrene-butadiene rubber, ethylene-ethyl acrylate copolymer, and ionomer.

25 Particularly useful propylene copolymers are those copolymers of propylene and one or more non-propylenic olefins. Propylene copolymers include random and block copolymers of propylene and an olefin selected from the group consisting of ethylene, C₄-C₁₀ 1-olefins, and C₄-C₁₀ dienes. Propylene copolymers also include random terpolymers of propylene and 1-olefins selected from the group consisting of ethylene and C₄-C₈ 1-olefins. In terpolymers having both ethylene and C₄-C₈ 1-olefins, the ethylene content is preferably 20 percent or less by weight. The C₄-C₁₀ 1-olefins include the linear and branched C₄-C₁₀ 1-olefins such as, for example, 1-butene, isobutylene, 1-pentene, 3-methyl-1-

butene, 1-hexene, 3,4-dimethyl-1-butene, 1-heptene, and 3-methyl-1-hexene. Examples of C₄-C₁₀ dienes include 1,3-butadiene, 1,4-pentadiene, isoprene, 1,5-hexadiene, 2,3-dimethyl-1,3-hexadiene.

5 Also, as used herein, the propylene polymer material has a melt flow rate of between 0.05 and 50 and preferably between 0.1 and 10 according to ASTM D1238 Condition L.

10 The preferred propylene polymer resins for the present invention are those polypropylene resins which are branched or lightly cross-linked polymer materials. Branching (or light cross-linking) may be obtained by those methods generally known in the art, such as
15 chemical or irradiation branching/light cross-linking. One such resin which is prepared as a branched/lightly cross-linked polypropylene resin prior to using the polypropylene resin to prepare a finished polypropylene
20 resin product and the method of preparing such a polypropylene resin is described in U.S. Patent No. 4,916,198, which is hereby incorporated by reference. Another method to prepare branched/lightly cross-linked polypropylene resin is to introduce chemical compounds
25 into the extruder, along with a polypropylene resin and allow the branching/lightly cross-linking reaction to take place in the extruder. U.S. Patent No. 4,714,716 illustrates this method and is incorporated by
30 reference.

 Suitable branching/crosslinking agents for use in extrusion reactions have been found to include azido and vinyl functional silanes, organic peroxides and multifunctional vinyl monomers.

It is also possible to add various additives such as inorganic fillers, pigments, antioxidants, acid scavengers, ultraviolet absorbers, flame retardants, processing aids, and extrusion aids.

5 In addition, a nucleating agent may be added in order to control the size of foam cells. Preferred nucleating agents include inorganic substances such as calcium carbonate, talc, clay, titanium oxide, silica,
10 barium sulfate, diatomaceous earth, and mixtures of citric acid and sodium bicarbonate. The amount of nucleating agent employed may range from 0.01 to 5 parts by weight per hundred parts by weight of a polymer resin. The preferred range is from 0.1 to 3 parts by
15 weight.

 Suitable inorganic blowing agents useful in making the foams of the present invention include carbon dioxide, nitrogen, argon, water, air, nitrogen, and
20 helium. The blowing agent will be comprised of at least 15 percent by weight, preferably at least 50 percent by weight, more preferably at least 95 percent by weight, and most preferably entirely of one or more of these inorganic agents. Preferred blowing agents include
25 carbon dioxide, nitrogen, and water. Most preferred blowing agents are carbon dioxide and a mixture of carbon dioxide and water. The balance of the blowing agent may comprise one or more chemical or volatile organic blowing agents. Preferred blowing agents,
30 however, are free of volatile organic agents and chemical blowing agents. Organic blowing agents include aliphatic hydrocarbons having 1-9 carbon atoms and halogenated aliphatic hydrocarbons, having 1-4 carbon atoms. Aliphatic hydrocarbons include methane, ethane, propane, n-butane, isobutane, n-pentane, isopentane,

neopentane. Among halogenated hydrocarbons, fluorinated hydrocarbons are preferred. Examples of fluorinated hydrocarbon include methyl fluoride, perfluoromethane, ethyl fluoride, 1,1-difluoroethane, 1,1,1-trifluoroethane (HFC-143a), 1,1,1,2-tetrafluoro-ethane (HFC-134a), pentafluoroethane, perfluoroethane, 2,2-difluoropropane, 1,1,1-trifluoropropane, perfluoropropane, perfluorobutane, perfluorocyclobutane. Partially halogenated chlorocarbons and chlorofluorocarbons for use in this invention include methyl chloride, methylene chloride, ethyl chloride, 1,1,1-trichloroethane, 1,1-dichloro-1-fluoroethane (HCFC-141b), 1-chloro-1,1-difluoroethane (HCFC-142b), 1,1-dichloro-2,2,2-trifluoroethane (HCFC-123) and 1-chloro-1,2,2,2-tetrafluoroethane (HCFC-124). Fully halogenated chlorofluorocarbons include trichloromonofluoromethane (CFC-11), dichlorodifluoromethane (CFC-12), trichlorotrifluoroethane (CFC-113), dichlorotetrafluoroethane (CFC-114), chloroheptafluoropropane, and dichlorohexafluoropropane. Fully halogenated chlorofluorocarbons are not preferred due to their ozone depletion potential. Chemical blowing agents include azodicarbonamide, azodiisobutyronitrile, benzenesulfonhydrazide, 4,4-oxybenzene sulfonyl-semicarbazide, p-toluene sulfonyl semicarbazide, barium azodicarboxylate, N,N'-dimethyl-N,N'-dinitrosoterephthalamide, and trihydrazino triazine.

The amount of blowing agent incorporated into the polymer melt to make a foam-forming polymer gel is from 0.2 to 5.0, preferably from 0.5 to 3.0, and most preferably from 1.0 to 2.50 moles per kilogram of polymer.

The present foam may be closed cell or open cell. Preferably, the present foam is from 5 to 100 and most preferably from 20 to 70 percent open cell according to ASTM D2856-A. Some open cell content is preferred in the present foam because an open cell foam is more dimensionally stable than a corresponding closed cell foam when prepared with fast permeating blowing agents such as carbon dioxide and water. The open cell foams prepared from the propylene polymer provide a high degree of resiliency most normally only seen in closed cell foams.

The present foam has the density of from 10 to 150 and most preferably from 10 to 70 kilograms per cubic meter. The foam has an average cell size of from 0.1 to 5.0 and preferably from 0.2 to 1.5 millimeters according to ASTM D3576.

The present foam has the cell wall thickness of less than 35, preferably less than 25, and most preferably less than 15 micrometers according to the formula below. It is believed that blowing agents diffuse out more rapidly from thinner cells walls than thicker ones providing more effective quenching and timely stiffening of the cell walls. As a result, the expanding bubbles become more effectively stabilized providing a low density foam having greater resiliency. The thickness of cell wall may be calculated approximately from the foam density and cell size by the use of following equation (1):

$$t = 0.46 D/B \quad (1)$$

where t is the thickness of cell walls in micrometers (μ): D is cell size in micrometers and: B, the

expansion ratio of the cellular body. Since the expansion ratio, B, can be equated the ratio of the polymer density (ρ_p) to foam density (ρ_f), the thickness of cell walls may be represented by equation (2):

5
$$t = (0.46/\rho_p)\rho_f D \quad (2)$$

For example, for polypropylene having 900 kg/m³ density to have cell wall thickness less than or equal to 35 micrometers:

10
$$(\rho_f)(D) \leq 68.5 \text{ (Kg/m}^3\cdot\text{mm)} \quad (3)$$

Equation 3 indicates that, for example, a 70 Kg/m³ density foam must have cell size no greater than 0.98mm. In order to meet the more preferred criterion $t \leq 15$ micrometers, a 70 kg/m³ density foam must have a cell size no greater than 0.42 mm.

The present foams may be made in any cross-sectional size or configuration such as foam sheet or plank, but are particularly useful in making plank foam having a major dimension in cross-section of at least 5 centimeters or a cross-sectional area of at least 10 square centimeters.

25 The present propylene polymer foam is generally prepared by heating a propylene polymer material to form a plasticized or melt polymer material, incorporating therein a blowing agent to form a foamable gel, and extruding the gel through a die to form the foam product. The blowing agent may be incorporated or mixed into the plastic melt by any means known in the art such as with an extruder, mixer, or blender. Prior to mixing with the blowing agent, the plastic material is heated to a temperature at or above the glass transition temperature or the melting point of the plastic

material. The blowing agent is mixed with the plastic melt at an elevated pressure sufficient to prevent substantial expansion of the melt and to generally disperse the blowing agent homogeneously within the melt. Optionally, a nucleator is blended in the polymer melt. The feeding rate of blowing agent and nucleator are adjusted to achieve a relatively low density foam and small cell size, which results in a foam having thin cell walls. After incorporation of the blowing agent, the foamable gel is typically cooled to a lower temperature to optimize physical characteristics of the foam product. The gel is then extruded through a die of desired shape to a zone of lower pressure to form the foam product.

Other suitable processes for making the present foam are the coalesced foam process as described in U.S. Patent No. 4,824,720 and the accumulating extrusion process described in U.S. Patent No. 4,323,528, both hereby incorporated by reference.

U.S. Patent No. 4,824,720, which describes the coalesced (or strand) foam process, is hereby incorporated by reference. This patent describes a method for providing a closed cell foam structure comprising a plurality of coalesced extruded strands or profiles by extrusion foaming of a molten thermoplastic composition utilizing a die containing a multiplicity of orifices. The orifices are so arranged such that the contact between adjacent streams of the molten extrudate occurs during the foaming process and the contacting surfaces adhere to one another with sufficient adhesion to result in a unitary structure. The individual strands of coalesced polyolefin foam should remain adhered into a unitary structure to prevent strand

delamination under stresses encountered in preparing, shaping, and using the foam.

U.S. Patent No. 4,323,528 describes an accumulating extrusion process and also is hereby
5 incorporated by reference. In this accumulating extrusion process low density, elongated cellular bodies having large lateral cross-sectional areas are prepared by: 1) forming, under pressure, a mixture of a thermoplastic polymer and a blowing agent, with the
10 mixture having a temperature at which the viscosity of the mixture is sufficient to retain the blowing agent when the mixture is allowed to expand; 2) extruding the mixture into a holding zone maintained at a temperature and pressure which does not allow the mixture to foam,
15 the holding zone having an outlet die defining an orifice opening into a zone of lower pressure at which the mixture foams, and an openable gate closing the die orifice; 3) periodically opening the gate; 4)
20 substantially concurrently applying mechanical pressure by a movable ram on the mixture to eject the mixture from the holding zone through the die orifice into the zone of lower pressure, at a rate greater than that at which substantial foaming in the die orifice occurs and
25 less than that at which substantial irregularities in cross-sectional area or shape occurs; and 5) permitting the ejected mixture to expand unrestrained in at least one dimension to produce an elongated thermoplastic
30 cellular body.

The present foams offer excellent cushioning properties and resiliency. The present foams are suitable for cushion packaging since the foams have adequate impact absorption properties and resilience. The foam products mitigate impact during multiple drops.

The more closed cell foams function better than the more open cell foams, but the open cell foams absorb dynamic impacts surprisingly well. In addition, the open cell foams can be used for sound absorption. The foams can be used for thermal insulation as well.

5

The present foams have satisfactory dimensional stability even though the foams are expanded with fast permeating blowing agents. Preferred foams recover ≥ 80 percent of initial volume within a month with initial volume being measured within 30 seconds after extrusion.

10

The following are examples of the present invention, and are not to be construed as limiting. Unless otherwise indicated, all percentages, parts, or proportions are by weight.

15

20

25

30

Example 1

An open-cell propylene copolymer foam was prepared according to the process of the present invention.

A Himont HMS (high melt strength) 2/98 ethylene/propylene random copolymer (polymer) with a melt flow rate (MFR) of 0.47 (ASTM D 1238 Condition L) was employed. Granular copolymer and an antioxidant concentrate were fed to an extruder at a total rate of approximately 113.4 kilograms/hour (kg/hr) (250 pounds/hour) (lbs/hr). The antioxidant concentrate contained 8.4 percent each of a hindered phenol-type antioxidant (Irganox 1010 by Ciba-Geigy Corp.) and a phosphite-type antioxidant (Ultranox 626 by General Electric Co.). The concentration of antioxidant in the extrudate was about 0.2 parts per hundred (pph) parts polymer.

The extruder was a 90 millimeter (3-1/2 inch) screw type having additional zone of mixing and cooling following the usual sequential zones of feeding, metering, and mixing. The temperatures maintained at the extruder zones were 130°C at the feeding zone, 200°C at the melting zone, 230°C at the metering zone and 210°C at the mixing zone. The extrudate was cooled to approximately 152°C in the cooling zone prior to foaming.

The blowing agents were injected into the same injection port into the extruder barrel between the metering and mixing zones. The blowing agents comprised carbon dioxide (CO₂) and water (H₂O) in amounts of 1.9

gram moles per kilogram of polymer (mpk) and 0.58 mpk, respectively.

A die having a gap-adjustable, rectangular orifice was situated after the cooling zone. The die width was fixed at 25.4 mm (1 inch). The die gap was readjusted to about 1.85 mm to prevent prefoaming. The die pressure at the critical die gap was 630 pounds per square inch gauge (psig) (4.3 megapascals (MPa)).

A stable, open-cell foam approximately 2.8 centimeters (cm) thick and 9.4 cm wide was produced. As shown in Table I, the foam exhibited excellent dimensional stability. The cell wall thickness is calculated to be about 11 micrometers.

Table I

Foam Density (kg/m ³) (1)	Cell Size (mm) (2)	Open Cell Content (3)	Dimensional Stability			
			Min. (4)	1 day (5)	1 week (5)	1 month (5)
31	0.69	64	95	96	105	105

Notes:

- (1) Density of foam body measured in about one month after foam expansion and expressed in kilograms per cubic meter (kg/m³).
- (2) Cell size in millimeters per ASTM D3576.
- (3) Open cell content in percentage per ASTM D2856-A
- (4) Minimum volume of foam body experienced during aging as a percentage of the initial volume determined within 30 seconds after foam expansion. The minimum occurred at 5 min. after extrusion for this foam.
- (5) Volume of foam body at the specified time after foam expansion as a percentage of the initial volume.

Example 2

An open-cell propylene copolymer foam in coalesced strand form was prepared according to the process of the present invention.

5

The apparatus employed was substantially the same as in Example 1 except that the die was replaced with a multi-orifice die. The multi-orifice die had a total of 450 holes of 0.041 inch (1.04 mm) in diameter arranged in an equilateral triangular pattern. The holes were each spaced-apart by 0.25 inch (6.35 mm), and form 15 rows of 30 holes.

10

The extrudate was of substantially the same composition as that made in Example 1, except that the feed rate of solids (polymer and antioxidant concentrate) to the extruder was lowered to 90.7 kgs/hr (200 lbs/hr). The feed rates of antioxidant concentrate, carbon dioxide, and water were lowered proportionately. The extrudate was cooled to 151°C in the cooling zone prior to foaming.

20

A stable foam body of coalesced strands with excellent strand-to-strand adhesion. At a die pressure of 4.1 MPa (600 psig), the foam showed no sign of prefoaming. The foam had an approximately rectangular cross-section of 5.1 cm in thickness and 18.0 cm in width. The foam had a density of 34.0 kg/m³ (2.12 pcf), cell size of 0.48 mm, and open cell content of 82 percent. The thickness of cell walls is calculated to be about 8.3 micrometers.

25

30

As shown in Table II, the foam exhibited excellent dimensional stability, and, for its high open content, a satisfactorily-high compressive strength to

render it suitable for cushion packaging and other applications. The foam also recovered well after high compression.

Table II

5	Dimensional Stability				Compressive Strength		
	Min. (1)	1 day	1 week (2)	1 month	V	E (3)	H
10	95	97	97	97	51	171	61

Notes:

- (1) Same as Table 1.
 (2) Same as Table 1.
 (3) Compressive strength at 25% deflection in kilopascals per ASTM D3575: V = vertical direction, E = extrusion direction, and H = horizontal direction

20 Example 3

An open-cell propylene homopolymer foam in coalesced strand form was prepared according to the present invention.

25

The apparatus employed was substantially the same as in Example 2.

Process conditions were substantially the same except as follows: a 90/10 blend of a Himont HMS polypropylene homopolymer having a melt flow rate of 0.42 and a 93.5/6.5 ethylene/acrylic acid (EAA) copolymer having a melt index (ASTM 1238 Condition E) of 5.5 was expanded with a CO₂/H₂O blowing agents at 1.5 mpk/0.57 mpk (6.6 pph/1.30 pph), and the solids

(polymers and antioxidant concentrate) were fed to the extruder at a rate of 136 kgs/hr (300 lbs/hr). The extrudate was cooled to 162°C in the cooling zone prior to foaming.

5 A stable foam body of coalesced strands with excellent strand-to-strand adhesion. At a die pressure of 4.8 MPa (700 psig), the foam showed no sign of prefoaming. The foam had an approximately rectangular cross-section of 6.4 cm in thickness and 19.3 cm in
10 width. The foam had a density of 39.4 kg/m³ (2.46 pcf), cell size of 1.29 mm, and open cell content of 69 percent. The thickness of the cell walls is calculated to be about 26 micrometers.

15 As shown in Table III, the foam exhibited excellent dimensional stability, and, for its high open content, a satisfactorily high compressive strength to render it suitable for cushion packaging and other
20 applications. The foam also recovered well after high compression.

Table III

25	Dimensional Stability				Compressive Strength		
	Min. (2)	1 day	1 week	1 month (3)	V	E	H (1)
	92	94	95	95	46	237	52

30

Notes:

- (1) Same as Table I except that the minimum occurred at 15 min. after extrusion.
- (2) Same as Table I
- (3) Same as Table II

Example 4

An open-cell propylene copolymer foam was prepared according to the present invention.

5 The apparatus employed was a 38 mm (1-1/2 inch) screw type extruder of substantially the same configuration as the extruder in Example 1 except as described below. The apparatus had a gap-adjustable die of 6.35 mm (0.25 in) width.

10 Foams were produced by the substantially the same procedure as in Example 1. Himont HMS 2/98 ethylene/propylene (E-P) random copolymer having a melt flow rate of 0.34 (ASTM D1238 Condition L) was employed.
15 The level of antioxidant (Irganox 1010 by Ciba-Geigy Corp.) was adjusted to 0.1 pph. The extruder was adjusted to a solids extrusion rate of 3.7 kgs/hr (8.1 lbs/hr). The temperatures in the extruder were 160°C at feeding zone, 200°C at melting zone, 200°C at metering
20 zone, and 190°C at mixing zone. A blowing agent or blowing agents were selected from carbon dioxide, water and nitrogen as represented in Table IVa. A predetermined amount of each blowing agent were injected
25 separately into the injection port.

 The temperature of the cooling zone was varied among the tests as necessary to provide a good foam. The foaming temperature was varied slightly between
30 152°C and 158°C as necessary. When the foaming temperature was reached, the die gap was readjusted to produce a foam of the largest cross-section possible free from prefoaming. The threshold die gap for prefoaming (called critical die gap hereinafter) was varied from about 0.36 mm to 1.1 mm. The die pressure

at the critical die gap ranged from 5.7 MPa (830 psig) to 12.1 MPa (1750 psig).

5 Foam samples of approximately 12 cm (5 in) in length were examined for processability, dimensional stability, and quality. The results are set forth in Tables IVa and IVb.

10 In Tests 4.1 through 4.4, 4.7 and 4.8, good foams having low densities, small cells, and satisfactory cross-sectional sizes were obtained using carbon dioxide, carbon dioxide/water, carbon dioxide/water/nitrogen, carbon dioxide/nitrogen, and water/nitrogen as blowing agents. The foams had 20-60 percent open cells, and exhibit excellent dimensional
15 stability.

20

25

30

Table IVa Processability and Quality

	Test No.	CO ₂ (1)	H ₂ O (2)	N ₂ (3)	Foam Size (4)	Foam Dens. (5)	Cell Size (6)	Wall Thick (7)	Open Cell (8)	Foam Qual. (9)
5	4.1	1.6	0	0	1.4	36	0.49	9	57	G
	4.2	2.0	0	0	1.2	36	0.32	7	60	SV
	4.3	1.5	0.5	0	0.9	44	0.37	8	40	G
	4.4	1.0	1.0	0	1.0	44	0.27	6	42	G
	4.5*	0.8	0.8	0	3.0	49	1.80	45	84	V
10	4.6*	0.5	1.5	0	3.4	44	3.24	73	61	H
	4.7	1.0	0.8	0.2	1.2	49	0.70	18	52	G
	4.8	1.3	0	0.3	0.6	44	0.17	4	20	G
	4.9*	0	2.0	0	2.7	192	5.40	530	55	C
	4.10	0	2.0	0.2	1.2	57	0.77	23	72	SC
15	4.11	0	1.6	0.3	0.9	48	0.31	8	44	G

Notes: *Not an example of this invention

(1), (2) and (3) gram-moles of carbon dioxide, water, and nitrogen mixed in per one kilogram of polymer

(4) Cross-sectional area of foam body in square centimeters

20 (5) Density of foam body aged for about a month in kilogram per cubic meters

(6) Cell size in millimeters determined per ASTM D3576

(7) Thickness of cell walls in micrometers

(8) Open cell content in percentage determined per ASTM 2856-A

(9) Quality of foam body;

25 G = good foam having uniform cell distribution

V = foam body contains a number of voids

SV = foam body contains some voids

H = foam body contains large holes

C = foam body severely collapses

SC = foam body experiences a slight collapse

30

Table IVb Dimensional Stability

Test No.	Minimum (1)	1 day (2)	1 week (2)	1 month (2)
5 4.1	100	104	102	103
4.8	87	99	101	101
4.11*	85	95	98	100

Notes:

10 (1) Minimum volume of foam body experienced during aging as a percentage of the initial volume which was determined within 30 seconds after foam expansion. The minimum occurred at 15 min. after extrusion for this foam.

15 (2) Volume of foam body at the specified time after foam expansion as a percentage of the initial volume.

*For this foam, the stability is expressed by foam volume as a percentage of the volume of one month-old foam body.

20

Example 5

Open-cell propylene copolymer foams were prepared by steady-state extrusion and by accumulating
25 extrusion according to the present invention.

Substantially the same apparatus and operating procedure of Example 4 were employed. The resin utilized was a Himont HMS 2/98 ethylene/propylene random
30 copolymer having a melt flow rate of 0.53 (ASTM D1238 Condition L). The temperatures maintained in the extruder were 170°C at feeding zone, 230°C at melting zone, 200°C at metering zone and 185°C at mixing zone. Carbon dioxide and mixtures of same with water were employed as the blowing agent.

The temperature of the cooling zone was adjusted among the tests to provide a good foam. The foaming temperature varied between 149 and 150°C. The critical die gap and die pressure ranged from about 0.65 mm to 0.71 mm and from 7.4 MPa (1070 psig) to 8.8 MPa (1270 psig).

In addition to data and samples taken at the critical die gap, larger samples of the same formulation were produced by an accumulating extrusion scheme using the adjustable die orifice (hereinafter referred to as pseudo-accumulation extrusion).

A pseudo-accumulating extrusion scheme was also utilized. After a steady extrusion sample was taken, the die orifice was closed to accumulate the foamable melt in the extrusion line until the die pressure increased by about 3.5 MPa (500 psi) and then the die was quickly opened to about 1.8-2.0 mm gap allowing the extrudate melt to exit the die and foam. The instantaneous extrusion rate was 8-9 kgs/hr (17-20 lbs/hr), about twice as high as the steady state extrusion rate. The pseudo-accumulating scheme simulates a high rate of steady extrusion. The pseudo-accumulating scheme produced good, lower-density foam strands free from corrugation. A fine-celled low-density foam often suffers corrugation of the foam body in the transverse direction when the extrusion rate is not sufficiently high enough. The foam samples produced by the pseudo-extrusion were used to determine dimensional stability. The processability, quality, and dimensional stability of the foams produced in this Example are set forth in Table Va and Vb, respectively.

The results of the tests demonstrate that relatively thin walls yield better quality foams, which are made at a higher level of carbon dioxide. A higher level of blowing agent not only provides a lower foam density but makes the cell size smaller. The foams produced in Tests 5.4a and 5.4b, with cell walls thicker than 35 microns, have large holes at the center. The foams made in Tests 5.2 and 5.2b have 30-31 micron cell walls and are marginally acceptable with some voids. Other foams having cell walls thinner than 10 microns are good in integrity with a uniform distribution of cell sizes. Two foams tested exhibit satisfactory dimensional stability. It was noted that the higher density foam made in Test 5.3b exhibits the better dimensional stability than the slightly lower density foam made in Test 5.1b.

20

25

30

Table Va Processability and Quality

	Test No.	CO ₂	H ₂ O	Foam Size	Foam Dens.	Cell Size	Wall Thick	Open Cell	Foam Qual.
	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(9)
5	5.1a	1.5	0.5	1.0	36	0.39	7	26	G
	5.1b			1.2	31	0.29	5	27	G
	5.2a	1.0	0.5	1.5	41	1.47	31	74	SV
	5.2b			2.1	38	1.55	30	71	SV
	5.3a	1.6	0	0.9	44	0.30	7	33	G
	5.3b			2.1	33	0.30	5	21	G
10	5.4a*	1.3	0	1.1	65	1.08	36	76	H
	5.4b*			2.1	71	1.01	37	52	H

Notes: *Not an example of this invention

- 15 (1) Test a's represent a steady state extrusion and b's a pseudo-accumulating extrusion.
 (2) and (3) G-moles of carbon dioxide and water mixed in per one kilogram of polymer
 (4) Cross-sectional area of foam body in square centimeters
 (5) Density of foam body aged for about a month in kilogram per cubic meters
 20 (6) Cell size in millimeters determined per ASTM D3576
 (7) Thickness of cell walls in micrometers
 (8) Open cell content in percentage determined per ASTM 2856-A
 (9) Quality of foam body;
 G = good foam having uniform cell distribution
 V = foam body contains a number of voids
 25 SV = foam body contains some voids
 H = foam body contains large holes

30

Table Vb Dimensional Stability

	Test No.	Minimum (1)	1 day (2)	1 week (2)	1 month (2)
5	5.1b	76	88	93	95
	5.3b	100	106	107	106

Notes:

- 10 (1) Minimum volume of foam body experienced during aging as a percentage of the initial volume which was determined within 30 seconds after foam expansion. The minimum occurred at 15 min. after extrusion for this foam.
- 15 (2) Volume of foam body at the specified time after foam expansion as a percentage of the initial volume.

Example 6

20 Open-cell propylene copolymer foams were prepared using a nucleating agent according to the present invention.

25 Substantially the same apparatus and operating procedure of Example 4 were employed. A higher melt flow rate (MFR) resin was foamed with relatively small amount of carbon dioxide blowing agent and a cell nucleating agent. The resin employed was a Himont HMS 2/98 ethylene/propylene random copolymer having a MFR of 7.0 (ASTM D1238 Condition L). The cell nucleating agent 30 was Hydrocerol CF-70 (Boehlinger Ingelheim KG). Hydrocerol CF-70 is a sodium carbonate/citric acid-type nucleating agent. A small amount (0.1 pph) Irganox 1010 antioxidant was incorporated.

The granular polypropylene resin was mixed with the antioxidant concentrate and 0.2 pph of nucleator

pellets and extruded at a uniform rate of 4.7 kgs/hr (10.4 lbs/hr). The temperatures maintained in the extruder were 190°C at feeding zone, 220°C at melting zone, 210°C at metering zone, and 180°C at mixing zone. Carbon dioxide was injected into the blowing agent injection port at a predetermined uniform rate. The temperature of the cooling zone was adjusted in each test to provide a good foam. The foaming temperature of the formulations was 138-139°C. The critical die gap and die pressure for the formulations in this example ranged from about 0.25 mm to 0.30 m and from 9.7 MPa (1410 psig) to 9.9 MPa (1430 psig).

As shown in Table VI, good foams having relatively high densities and uniformly small cell sizes were obtained using 0.6-0.9 mpk (2.6-4.0 pph) carbon dioxide. The good processability of the foams may in part be due to their relatively thin cell walls of 11-15 microns. Even though the foams had relatively low open cell contents, the foams were not observed to suffer unsatisfactory shrinkage during aging (not actually measured). The high compressive strengths of the high density foams are believed to have aided in the dimensional stability of those foams.

30

Table VI Processability and Quality

Test No.	CO ₂ (1)	Foam Size (2)	Foam Dens. (3)	Cell Size (4)	Wall Thick (5)	Open Cell (6)	Foam Qual. (7)
5							
6.1	0.9	0.6	53	0.44	12	17	G
6.2	0.8	0.6	59	0.36	11	7	G
6.3	0.6	0.5	70	0.42	15	16	G

Notes: *Not an example of this invention

- 10 (1) gram-moles of carbon dioxide and water mixed in per
one kilogram of polymer
(2) Cross-sectional area of foam body in square
centimeters
(3) Density of foam body aged for about a month in
kilogram per cubic meters
(4) Cell size in millimeters determined per ASTM D3576
15 (5) Thickness of cell walls in micrometers
(6) Open cell content in percentage determined per ASTM
2856-A
(7) Quality of foam body;
G = good foam having uniform cell distribution

20

Example 7

25 Open-cell propylene copolymer foams with some
ethylenic polymer content were prepared according to the
present invention.

30 The apparatus and its operating procedure used
in this example were substantially the same as in
Example 5. The PP copolymer resin of Example 5 was
blended with 10-20 percent of various ethylenic polymer
resins and foamed with carbon dioxide or its mixture
with water.

The ethylenic resins evaluated included
Primacore 1410 ethylene/acrylic acid (EAA) copolymer (9

percent AA, 1.5 melt index) of The Dow Chemical Co., PE 682 low density polyethylene (LDPE) (0.7 melt index) of The Dow Chemical Co., and Surlyn 1706 brand ionomer (Ionomer) by E.I. DuPont de Nemours & Co. A small amount (0.1 pph) Irganox 1010 antioxidant was
5 incorporated in the polymers.

The granular PP copolymer resin and a predetermined amount of a selected ethylenic resin were dry-blended with the antioxidant concentrate. The solid
10 was extruded at a uniform rate of 3.7 kgs/hr (8.1 lbs/hr). The extruder zone temperatures were the same as in Example 5. The temperature of the cooling zone was adjusted in each of the tests of this example to
15 provide a good foam. The foaming temperature of the formulations was 147-149°C. Foams were made by conventional steady extrusion and pseudo-accumulating extrusion. The critical die gap and die pressures ranged from about 0.69 mm to 0.71 mm and from 8.6 MPa
20 (1250 psig) to 9.1 MPa (1320 psig).

The processability, quality, and dimensional stability data is set forth in Tables VIIa and VIIb. All blends of the PP resin and ethylenic polymer yielded
25 good quality foams having fine cells and low densities. The cell walls of the test foams are all thinner than 24 microns. It was surprising that polypropylene can accommodate as much as 20 percent of those ethylenic copolymers in expanding good low-density foams. Except
30 for the foam made in Test 7.4b, the foams displayed satisfactory dimensional stability with open cell content greater than 30 percent. The PP/ionomer blend foam made in Test 7.4b exhibited marginal dimensional stability with excessive shrinkage (more than 50 percent) and slow recovery. Desirable foams recover at

least 80 percent of its original volume within about a month. The low density foam of low open cell content was marginal in dimensional stability.

5

10

15

20

25

30

Table VIIa Processability

Test No.	Blend Resin		CO ₂	H ₂ O	Foam Size	Foam Dens.	Cell Size	Wall Thick	Open Cell	Foam Qual.
	Type	level								
(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)
7.1a	EAA	10	1.7	0	1.3	30	0.54	8	31	G
7.1b					2.9	31	0.52	8	34	G
7.2a	EAA	10	1.5	0.5	1.7	33	0.68	11	59	G
7.2b					5.0	30	0.62	10	50	G
7.3a	LDPE	20	1.5	0.5	1.1	37	0.34	6	39	G
7.3b					1.6	38	0.51	10	34	G
7.4a	IONOMER	10	1.5	0.5	ND	ND	ND	ND	ND	G
7.4b					4.6	29	0.52	8	10	G

Notes: ND = Not determined

(1) Test a's represent a steady state extrusion and b's a pseudo-accumulating extrusion

(2) The type of ethylenic polymer blended in with the PP copolymer:

20 EAA = ethylene acrylic acid copolymer (9 percent AA, 1.5 melt index), Primacore 1410 resin by Dow Chemical Co.

LDPE = Dow low density polyethylene 682 (0.7 melt index)

Ionomer = Surlyn 1706 grade ionomer made by DuPont de Nemours & Co.

(3) The percentage of the ethylenic polymer blended in the total polymer

25 (4) and (5) G-moles of carbon dioxide and water mixed in per one kilogram of polymer

(6) Cross-sectional area of foam body in square centimeters

(7) Density of foam body aged for about a month in kilogram per cubic meters

(8) Cell size in millimeters determined per ASTM D3576

(9) Thickness of cell walls in micrometers

30 (10) Open cell content in percentage determined per ASTM 2856-A

(11) Quality of foam body;

G = good foam having uniform cell distribution

Table VIIb Dimensional Stability

5	Test No.	Minimum (1)	1 day (2)	1 week (2)	1 month (2)
	7.1b	91	95	98	99
	7.2b	93	95	98	98
	7.3b	82	89	82	82
	7.4b	49	49	67	78

Notes:

- 10 (1) See Table Vb.
(2) See Table Vb.

15 Example 8

Open-cell propylene copolymer foams were prepared according to the present invention.

20 The apparatus employed was a 25 mm (1 inch) screw type extruder of substantially the same configuration as that of Example 4. The adjustable die gap was 3.68 mm (0.145 inch) in width.

25 The operating procedure was substantially the same as in Example 4. The same E-P copolymer of Example 4 was dry-blended with EAA resin, Ionomer resin, and 0.1 pph Irganox 1010, and fed to the extruder at a rate of 1.8 kgs/hr (4 lbs/hr). A binary blowing agent
30 consisting of 0.8 mpk CO₂ and 0.8 mpk H₂O was injected into the extruder barrel. The temperatures maintained at the extruder zones were 160°C at feeding zone, 180°C at melting zone, 210°C at metering zone and 230°C at mixing zone. Carbon dioxide and water were separately injected into the injection port each at a rate to be

0.8 mpk. The temperature of the cooling zone was adjusted in each test to provide a good foam. The foaming temperatures of the formulations evaluated in this example ranged from 165°C to 168°C. The critical die gaps and the critical pressure were in the range of
5 0.36-0.51 mm (14-20 mils) and 4.5-7.9 MPa (650-1150 psig), respectively.

The processability, quality, and dimensional stability of the foams are set forth in Tables VIIIA and
10 VIIIB. Both the E-P copolymer and its blends provided satisfactory foams having low foam densities and acceptable dimensional stabilities. While the foams having cell walls of 10 micron thickness were of
15 excellent quality, those having 21-22 micron-thick cell walls were marginally satisfactory in quality. Tests 8.1, 8.2 and 8.3 indicate that addition of a small amount of EAA resin aids in foam expansion of the E-P copolymer with the relatively large amount of water
20 blowing agent. The EAA resin may enhance solubility of water in the polymer melt.

25

30

Table VIIa Processability

Test No.	Blend Resin		CO ₂	H ₂ O	Foam Size	Foam Dens.	Cell Size	Wall Thick	Open Cell	Foam Qual.
	Type level									
	(1)	(2)								
8.1	-	-	0.8	0.8	0.7	44	0.95	21	81	D
8.2	EAA	5	0.8	0.8	0.7	40	0.51	10	21	G
8.3	EAA	10	0.8	0.8	1.0	35	1.25	22	83	SD
8.4	Ionomer	5	0.8	0.8	0.6	41	0.49	10	32	G

Notes:

(1) The type of ethylenic polymer blended in with the PP copolymer:

(2) The percentage of the ethylenic polymer blended in the total polymer

(3) and (4) G-moles of carbon dioxide and water mixed in per one kilogram of polymer

(5) Cross-sectional area of foam body in square centimeters

(6) Density of foam body aged for about a month in kilogram per cubic meters

(7) Cell size in millimeters determined per ASTM D3576

(8) Thickness of cell walls in micrometers

(9) Open cell content in percentage determined per ASTM 2856-A

(10) Quality of foam body;

G = good foam having uniform cell distribution

D = distribution of cell size

SD = slight distribution of cell size

Table VIIb Dimensional Stability

Test No.	Minimum (1)	1 day (2)	1 week (2)	1 month (2)
8.1	92	92	93	93
8.2	67	83	93	108
8.3	102	103	102	103
8.4	(not determined)			

Notes:

(1) See Table Vb

(2) See Table Vb

Example 9

Open-cell propylene copolymer foams were prepared by an accumulating extrusion system according to the present invention.

5

The apparatus consisted of the 38 mm (1-1/2 inch) foam extruder of Example 4, an accumulating extrusion system (AES) attached at the end of the cooling zone of the extruder, and a gap-adjustable die having an opening of 6.35 mm (0.25 in) width attached therewith. The foamable melt is accumulated in the accumulating chamber of 330 cubic centimeter capacity and then rapidly pushed out of the orifice by nitrogen pressure.

15

In this example, a 90/10 blend by weight of a 2/98 E-P copolymer (0.52 MFR) and Primacore® 1410 EAA copolymer was expanded with a binary blowing agent consisting of 1.5 mpk carbon dioxide and 0.5 mpk water. The solids were pre-blended, and extruded at a rate of 3.7 kgs/hr (8.1 lbs/hr). The temperatures in the extruder were 170°C at feeding zone, 200°C at melting zone, 220°C at metering zone, and 180°C at mixing zone. Carbon dioxide and water were separately injected into the injection port at a rate to be 1.5 and 0.5 mpk, respectively. The temperature of the cooling zone was adjusted to provide a good foam. The foaming temperature of the formulation was approximately 150°C. The critical die gap and die pressure were 0.8 mm and 8.0 MP (1160 psig), respectively.

20

25

30

As shown in Table IXa, a good foam having fine cell size was obtained by steady extrusion. The foam

strand produced had some corrugation in the transverse direction.

The AES scheme was employed to produce larger foam than possible with steady extrusion. The
5 temperatures of the accumulator and the transfer lines were maintained so that the foamable melt coming out of the extruder could be neither excessively cooled nor heated. The AES was operated as follows. First, the
10 chamber was pressurized with nitrogen to about 700 psig, which was to exert back pressure for the incoming melt. Then, the extruder output was diverted into the accumulator. The accumulation lasted for approximately 3.5 minutes. The gas pressure increased and exceeded
15 8.3 MPa (1200 psig) as it was compressed by the incoming melt. During the accumulating operation, the die gap was opened to 3.1 mm (120 mils), and the nitrogen source pressure was adjusted to 9.0 MPa (1300 psig). Upon
20 completion of accumulation, the accumulated melt was allowed to shoot out of the die orifice.

An excellent-quality foam free from corrugation was achieved. Several such shots were made to produce
25 foams for property evaluation. The instantaneous extrusion rates, estimated from the weights of the extrudates and the shoot-out times, were ranged from 90 to 120 kgs/hr. The foams were observed to be dimensionally stable.

30 The foam plank produced by the AES scheme was aged for a month, and tested for its physical properties. The data is summarized in Table IXb. After the skins were taken off, the density of the AES foam was 24 kg/m^3 (1.5 pcf), much lower than that with skin on (refer to Table IXa). The low-density foam having

over 60 percent open cells displays strength,
resiliency, and shock-mitigating property suitable for
cushion packaging of relatively light weight items.

5

10

15

20

25

30

Table IXa Processability and Quality

Test No.	Blend Resin		CO ₂	H ₂ O	Foam Size	Foam Dens.	Cell Size	Wall Thick	Open Cell	Foam Qual.
	Type level									
	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(9)	(10)
9.1a	EAA	10	1.5	0.5	1.6	39	0.58	12	36	G
9.1b					8.9	30	0.62	10	67	G

Notes:

- (1) Type of resin employed
 (2) The percentage of the EAA copolymer blended in the total polymer
 (3) and (4) gram-moles of carbon dioxide and water mixed in per one kilogram of polymer
 (5) Cross-sectional area of foam body in square centimeters
 (6) Density of foam body aged for about a month in kilogram per cubic meters
 (7) Cell size in millimeters determined per ASTM D3576
 (8) Thickness of cell walls in micrometers
 (9) Open cell content in percentage determined per ASTM 2856-A
 (10) Quality of foam body;
 G = good foam having uniform cell distribution

Table IXb Properties

	Property	Unit	Method	Condition	Value
5	Compressive Strength	kPa	ASTM D3575B	@ 10% defl.	38
				25% defl.	47
				50% defl.	65
	Compressive Recovery	percent	Dow (1)	after 1 hr	86
				after 6 days	93
10	Dynamic Cushioning	G	Dow (2)	@ static stress (kPa)	
				1.8	77
				3.5	70
				5.2	75
				8.6	104
15	Recovery After D.C.	percent	Dow (3)	@ static stress (kPa)	
				1.8	95
				3.5	98
				5.2	98
				8.6	97

20 Notes: *All properties are for the vertical direction

- (1) Thickness of the specimen in percentage of the initial six days after it was compressed to 80 percent of the initial at rate of 1.27 cm/min (0.5 inch/min.)
- (2) Average of peak deceleration during second to fifth drops in g's at the given static stress. The test specimens had 38 mm (1.5 inches) thickness and 50 mm x 50 mm (2 inches x 2 inches) impact area. The drop height of the weights was 61 cm (24 inches).
- (3) Thickness of the specimen in percentage of the initial six days after is was tested for the dynamic cushioning property.

30 While embodiments of the propylene polymer foam have been shown with regard to specific details, it will be appreciated that depending upon the properties of the foam and the manufacturer's desires the present invention may be modified by various changes while still being fairly within the scope of the novel teachings and principles herein set forth.

CLAIMS

1. The process for making a dimensionally-stable, extruded propylene polymer foam having a density of from 10 to 150 kilograms per cubic meter and an average cell wall thickness of less than 35 micrometers,
5 comprising:
 - a) heating a propylene polymer material comprising greater than 50 percent by weight propylene monomeric units to form a melt polymer material;
 - 10 b) incorporating into the melt polymer material a blowing agent to form a foamable gel;
 - c) cooling the foamable gel to an optimum foaming temperature; and
 - 15 d) extruding the foamable gel through a die to form the foam, the process being characterized in that the blowing agent contains at least 15 percent by weight of one or more inorganic blowing agents.
2. The process of Claim 1, wherein the blowing
20 agent contains carbon dioxide.
3. The process of Claim 1, wherein the blowing agent contains carbon dioxide and water.

4. The process of Claim 1, wherein the blowing agent contains nitrogen and water.

5. The process of any of Claims 1 through 4, wherein the blowing agent incorporated into the melt polymer material comprises at least 50 weight percent by weight of one or more inorganic blowing agents.

6. The process of any of Claims 1 through 4, wherein the blowing agent incorporated into the melt polymer material is at least 95 weight percent by weight of one or more inorganic blowing agents based on the total weight of the blowing agent.

7. The process of any of Claims 1 through 4, wherein the blowing agent incorporated into the melt polymer material is entirely of one or more inorganic blowing agents.

8. The process of any of Claims 1 through 4, wherein the blowing agent is incorporated into the melt polymer material at a concentration of from 0.2 to 5.0 moles per kilogram of melt polymer material.

9. A foam obtainable from the process of any of Claims 1 through 4.

30

INTERNATIONAL SEARCH REPORT

L national application No.
PCT/US92/10932

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) :C08J 9/08, 9/10, 9/12

US CL :264/51; 521/82, 94, 97, 142, 143

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 264/51; 521/82, 94, 97, 142, 143

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US,A, 3,407,151 (OVERCASHIER et al) 22 October 1968 Entire document.	1-24
A	US,A, 4,506,037 (SUZUKI et al) 19 March 1985 Entire document.	1-24

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be part of particular relevance	*X*	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
E earlier document published on or after the international filing date	*Y*	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z*	document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means		
P document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

11 MARCH 1993

Date of mailing of the international search report

26 APR 1993

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. NOT APPLICABLE

Authorized officer

MORTON FOELAK

Telephone No. (703) 308-2351